Reply to Office Action of September 4, 2008

AMENDMENTS TO THE CLAIMS

Docket No.: ALXN-P02-089

This listing of the claims will replace all prior versions, and listings, of claims in the

application:

1. (Currently Amended) A method for reducing an immune response in an animal in

need thereof by inhibiting an interaction between a dendritic cell and a T cell, comprising

administering to an animal in need of reducing said immune response a compound an antibody

which binds to a protein with the amino acid sequence of SEQ ID NO: 2 (DC-SIGN) on the surface

of a dendritic cell, wherein said antibody compound-reduces one or more interactions between a

dendritic cell and a T cell thereby reducing said immune response in the animal, and wherein the

animal is not infected with HIV.

2. (Canceled)

3. (Original) The method of claim 1 wherein said animal is a mammal.

4. (Original) The method of claim 3 wherein said mammal is a human.

5. (Canceled)

6. (Currently amended) The method of claim 1 wherein said antibody compound

reduces adhesion between DC-SIGN and an ICAM receptor on the surface of a T cell.

7. (Original) The method of claim 6 wherein said ICAM receptor is selected from the

group consisting of ICAM-2 receptors and ICAM-3 receptors.

Claims 8 - 18. (Canceled)

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19. (Currently amended) The method of claim [[9]] 1 wherein said antibody is a monoclonal antibody.

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Claims 20 - 22. (Canceled)

- 23. (Currently amended) The method of claim [[9]] 1 wherein said antibody is selected from the group consisting of i) an antibody produced by hybridoma ECACC accession number 99040818 and ii) an antibody produced by hybridoma ECACC accession number 99040819.
- 24. (Previously Presented) The method of claim 1, wherein said animal is in need of tolerance, immunotherapy or immunosuppression.
- 25. (Previously Presented) The method of claim 1, wherein said animal is suffering from an autoimmune disease.
- 26. (Previously Presented) The method of claim 1, wherein said animal is suffering from an allergy.
- 27. (New) The method of claim 1, wherein the antibody is administered in combination with another compound selected from the group consisting of: immunosuppressants, immunomodulants, antibiotics, auto-antigens, allergens, anti-LF3A, Tumor Necrosis Factor (TNF), anti-viral agents, and CD4 inhibitors.